# NEOSPORIN ORIGINAL- bacitracin zinc, neomycin sulfate, and polymyxin b sulfate ointment Johnson & Johnson Consumer Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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#### **NEOSPORIN ORIGINAL OINTMENT**

## **Drug Facts**

Active ingredients (in each gram)	Purpose
Bacitracin 400 units	First aid antibiotic
Neomycin 3.5 mg	First aid antibiotic
Polymyxin B 5,000 units	First aid antibiotic

#### Use

first aid to help prevent infection in minor:

- cuts
- scrapes
- burns

## **Warnings**

# For external use only.

#### Do not use

- if you are allergic to any of the ingredients
- in the eyes
- over large areas of the body

# Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

# Stop use and ask a doctor if

- you need to use longer than 1 week
- condition persists or gets worse
- rash or other allergic reaction develops

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

- clean the affected area
- apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
- may be covered with a sterile bandage

### Other information

• store at 20° to 25°C (68° to 77° F)

## **Inactive ingredients**

cocoa butter, cottonseed oil, olive oil, sodium pyruvate, vitamin E, white petrolatum

# **Questions?**

call **1-800-223-0182** 

# PRINCIPAL DISPLAY PANEL - 14.2 g Carton ORIGINAL OINTMENT NEOSPORIN®

Pacitrasin Zine Noomyo

Bacitracin Zinc-Neomycin Sulfate-Polymyxin B Sulfate FIRST AID ANTIBIOTIC OINTMENT

NET WT 0.5 oz (14.2 g)



### **NEOSPORIN ORIGINAL**

bacitracin zinc, neomycin sulfate, and polymyxin b sulfate ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58232-4002
Route of Administration	TOPICAL		

## **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I)	BACITRACIN	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII: J2VZ 07J96K)	POLYMYXIN B	5000 [iU] in 1 g

Inactive Ingredients			
Ingredient Name	Strength		
COCOA BUTTER (UNII: 5120YT1CRR)			
COTTONSEED OIL (UNII: H3E878020N)			
OLIVE OIL (UNII: 6UYK2W1W1E)			
SODIUM PYRUVATE (UNII: POD38AIF08)			
.ALPHATOCOPHEROL (UNII: H4N855PNZ1)			
PETROLATUM (UNII: 4T6H12BN9U)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58232- 4002-1	0.9 g in 1 POUCH; Type 0: Not a Combination Product	08/03/2011	06/14/2018
2	NDC:58232- 4002-2	1 in 1 CARTON	08/03/2011	06/14/2018
2		14.2 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:58232- 4002-6	1 in 1 CARTON	08/03/2011	07/18/2018
3		28.3 g in 1 TUBE; Type 0: Not a Combination Product		
4	NDC:58232- 4002-9	144 in 1 CARTON	08/03/2011	
4		0.9 g in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:58232- 4002-8	0.9 g in 1 POUCH; Type 0: Not a Combination Product	08/03/2011	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	08/03/2011	

Labeler - Johnson & Johnson Consumer Inc. (002347102)